

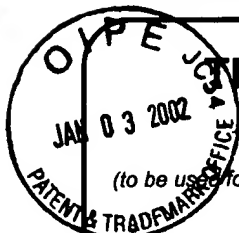
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PTO/SB/21 (08-00)

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TRANSMITTAL FORM

(to be used for all correspondence after initial filing)

Applicati n Number	09/932,538
Filing Date	August 17, 2001
First Nam d Inventor	Bell
Group Art Unit	1645
Examiner Name	
Attorney Docket Number	37070/205236

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ENCLOSURES (check all that apply)

☐ Fee Transmittal Form

☐ Fee Attached

☒ Preliminary Amendment

☐ After Final

☐ Affidavits/declaration(s)

☐ Extension of Time Request

☐ Express Abandonment Request

☐ Information Disclosure Statement

☐ Certified Copy of Priority Document(s)

☐ Response to Missing Parts/ Incomplete Application

☐ Response to Missing Parts under 37 CFR 1.52 or 1.53

☐ Assignment Papers (for an Application)

☒ Drawing(s)

☐ Licensing-related Papers

☐ Petition

☐ Petition to Convert to a Provisional Application

☐ Power of Attorney, Revocation Change of Correspondence Address

☐ Terminal Disclaimer

☐ Request for Refund

☐ CD, Number of CD(s) _____

☐ After Allowance Communication to Group

☐ Appeal Communication to Board of Appeals and Interferences

☐ Appeal Communication to Group (Appeal Notice, Brief, Reply Brief)

☐ Proprietary Information

☐ Status Letter

☐ Other Enclosure(s) (please identify below):

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Remarks

SIGNATURE OF APPLICANT, ATTORNEY, OR AGENT

Firm or Individual name

Kristin D. Mallatt, Reg. No. 46,895

Signature

Kristin D. Mallatt

Date

November 16, 2001

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APPLICANTS: Bell, et al.

SERIAL NO.: 09/932,538

GROUP ART UNIT:

FILED: August 17, 2001

EXAMINER:

FOR: CALCIUM PHOSPHATE PARTICLES AS MUCOSAL ADJUVANTS

Attorney Docket No.: 37070/205236

Commissioner for Patents
Washington, D.C. 20231

DATE: November 16, 2001

4/A
JM
4/12/02

PRELIMINARY AMENDMENT

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Sir:

Prior to any examination, please preliminarily amend the above-identified application as indicated below:

In the Specification

Kindly amend the specification as follows:

On page 33, line 26, before the sentence beginning "Serological" please insert the following paragraph:

Figures 4a and 4b show results from BALB/c mice (n=5/group) immunized on days 0 and 7 by intranasal or intravaginal delivery of 20 µg HSV-2 proteins alone, 51 µg CAP alone or HSV-2+ CAP (20 µg protein and 51 µg (CAP). Samples were

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